

USE OF COMPOUNDS THAT REDUCE ALPHA2-ANTIPLASMIN IN VIVO FOR THE  
PREPARATION OF A COMPOSITION FOR THE TREATMENT OF ISCHEMIC STROKE

*the following is a summary of the invention*

The present invention relates to a new means  
for the treatment of focal ischemic cerebral infarction  
5 (ischemic stroke).

Focal ischemic cerebral infarction occurs when  
the arterial blood flow to a specific region of the brain  
is reduced below a critical level resulting in neuronal  
cell death. It is thought that neuronal degeneration in  
10 central nervous system (CNS) diseases such as stroke,  
epilepsy and Alzheimer's disease is stimulated by an  
excess of the excitatory amino acid glutamate (2).  
Injection of glutamate agonists in the CNS indeed induces  
hippocampal neuronal cell death similar to that observed  
15 in neurodegenerative diseases (3).

Excitotoxin-induced neuronal degeneration is  
mediated by tissue-type plasminogen activator (t-PA) (4).  
Consistent with this observation, mice deficient in t-PA  
are resistant to, and infusion of plasminogen activator  
20 inhibitor-1 (PAI-1) protects against excitotoxin-mediated  
hippocampal neuronal degeneration (4-6).

Furthermore, deficiency of plasminogen (Plg),  
the zymogen substrate of t-PA, and infusion of  
 $\alpha_2$ -antiplasmin ( $\alpha_2$ -AP), protect mice against  
25 excitotoxin-induced hippocampal neuronal death (5). It  
has been proposed that plasmin-mediated degradation of  
laminin sensitizes hippocampal neurons to cell death by  
disrupting neuron-extracellular matrix interaction (7).

Wang et al. (8) recently demonstrated that  
30 neuronal damage after focal cerebral ischemia induced by  
transient occlusion of the middle cerebral artery was  
also reduced in mice with t-PA deficiency and exacerbated  
by t-PA infusion. This suggests that the plasminogen  
system may be involved both in establishing a cerebral  
35 ischemic infarct and in its extension during thrombolytic  
therapy. It was recently demonstrated that the neurotoxic  
effect of t-PA on persistent focal cerebral ischemia also  
occurred with other thrombolytic agents, including

streptokinase and staphylokinase (9). Thus, in those patients with persistent cerebral arterial occlusion, thrombolytic therapy for ischemic stroke may cause infarct extension, which would not only partially offset the established overall beneficial effect of arterial recanalization (10, 11), but indeed be harmful to a subgroup of patients. Because it is not possible to distinguish between patients who will and those who will not achieve cerebral arterial recanalization with thrombolytic therapy, the development of specific conjunctive strategies to counteract the neurotoxic effects of thrombolytic agents on persisting focal cerebral ischemia appear to be warranted.

It is therefore the object of the present invention to provide a new means for treating ischemic stroke.

In the research that led to the present invention the following was contemplated. Although it is assumed that neuronal injury during focal ischemia in the brain occurs primarily as a result of accumulation of excitotoxins such as glutamates, the role of plasmin-mediated laminin degradation or alternative mechanisms in the pathogenesis of cortical neuronal cell death has not been demonstrated. In order to delineate the contribution of individual components of the plasminogen (fibrinolytic) system on focal cerebral ischemic infarction, the present inventors then quantitated infarct size produced by ligation of the left middle cerebral artery (MCA) in mice with targeted inactivation of the genes encoding Plg, its activators tissue-type plasminogen activator (t-PA) or urokinase-type plasminogen activator (u-PA), or the fibrinolytic inhibitors PAI-1 or  $\alpha_2$ -AP. In addition, the effects of adenoviral transfer of the t-PA and PAI-1 genes and of infusion of human  $\alpha_2$ -AP on cerebral infarction were studied.

Whereas the findings of Strickland et al., that t-PA deficiency protects against focal cerebral ischemic

infarction were fully confirmed, and extended by the observation that PAI-1 deficiency resulted in significantly larger infarct sizes, the observation that Plg deficiency protects against excitotoxin induced neuronal cell death could not be confirmed. Instead it was found that focal cerebral infarct size was significantly larger in mice with Plg deficiency and conversely, significantly smaller in mice with  $\alpha_2$ -AP deficiency.

In aggregate, these findings indicate that plasminogen system components affect focal cerebral ischemic infarct size at two different levels: 1) reduction of t-PA activity (t-PA gene inactivation or PAI-1 gene transfer) reduces, while its augmentation (t-PA gene transfer or PAI-1 gene inactivation) increases infarct size, and 2) reduction of Plg activity (Plg gene inactivation or  $\alpha_2$ -AP injection) increases, while its augmentation ( $\alpha_2$ -AP gene inactivation or  $\alpha_2$ -AP neutralization) reduces infarct size. The findings are incompatible with a unique linked pathway in which t-PA-mediated plasmin generation would lead to neuronal cell death, but suggests two independent (t-PA mediated and Plg-mediated, respectively) mechanisms operating in opposite direction.

The internally consistent observations with  $\alpha_2$ -AP were unexpected but are most relevant for the treatment of ischemic stroke. Firstly a correlation was found between infarct size and genotype with heterozygotes displaying infarct sizes between those of the wild type and homozygous phenotypes. Secondly, bolus injection of human  $\alpha_2$ -AP (h $\alpha_2$ -AP) in  $\alpha_2$ -AP<sup>-/-</sup> mice caused a dose-related infarct expansion. Importantly, Fab fragments from affino-specific polyclonal rabbit anti-h $\alpha_2$ -AP antibodies, given intravenously 40 min after occlusion of the MCA, significantly reduced the cerebral ischemic infarct size. This observation suggests a potential avenue to counteract focal ischemic infarction with the use of  $\alpha_2$ -AP inhibitors (e.g. neutralizing monoclonal antibodies or compounds neutralizing  $\alpha_2$ -AP

activity). This approach was confirmed by infusion of plasmin in mice with MCA occlusion which, by neutralizing  $\alpha_2$ -AP, significantly reduced infarct size. The concentration of  $\alpha_2$ -AP in human plasma is 1 mM (12),  
5 corresponding to a total body pool of approximately 500 mg. An equivalent dose of a monoclonal Fab fragment would be approximately 400 mg, and that of plasmin approximately 500 mg, which is high but not excessive for single therapeutic administration. Furthermore, the  
10 observation that infarct size is proportional to the  $\alpha_2$ -AP level (derived from the gene dose effect and the dose-response) suggests that a partial reduction of the plasma level might have a significant beneficial effect.

In view of the above the invention thus relates  
15 to the use of compounds that reduce  $\alpha_2$ -AP activity in vivo for the treatment of focal cerebral ischemic infarction (ischemic stroke).

In a specific embodiment of the invention use is made of compounds that reduce the circulating  $\alpha_2$ -AP  
20 concentration. A lower concentration of  $\alpha_2$ -AP will lead to a lower activity. In an alternative embodiment, the activity of circulating  $\alpha_2$ -AP is reduced directly.

Compounds that are suitable for the reduction of  $\alpha_2$ -AP concentration and activity are for example  $\alpha_2$ -AP  
25 neutralizing antibodies or derivatives thereof. Preferred antibodies are monoclonal antibodies. Derivatives are preferably Fab fragments, scFv fragments.

Compounds neutralizing  $\alpha_2$ -AP are for example plasmin, mini-plasmin (lacking the first 4 kringles) or  
30 micro-plasmin (lacking all five kringles).

The present invention will be demonstrated in more detail in the following examples, which are however not intended to be limiting to the scope of the invention. In the examples reference is made to the  
35 following drawings:

Figures 1 to 3 are histograms comparing the volume (in mm<sup>3</sup>) of focal cerebral ischemic infarcts after ligation of the middle cerebral artery (MCA) in mice. The

data represent mean values and the vertical bars SEM, with the number of experiments given in the columns.

Figure 1 shows the effect of deficiency of plasminogen system components (genotype in abscissa) on focal ischemic cerebral infarct size (in mm<sup>3</sup>). WT: wild type (pooled values of 50% C57BL6/50% S129, 100% C57BL6 and 100% S129 genetic background).

Figure 2 shows the effect of adenoviral transfer of the t-PA or PAI-1 genes on focal ischemic cerebral infarct size in t-PA or PAI-1 deficient mice, respectively.

Figure 3 shows the effect of  $\alpha_2$ -AP on focal ischemic cerebral infarct size.

A. Effect of  $\alpha_2$ -AP genotype on cerebral infarct size.

B. Effect of injection of h $\alpha_2$ -AP or of h $\alpha_2$ -AP followed by anti-h $\alpha_2$ -AP Fab fragments on cerebral infarct size.

## 20 EXAMPLES

### EXAMPLE 1

#### Murine cerebral ischemic infarction model

##### 1. Introduction

All mice included in the present study were generated and bred at the Specific Pathogen Free Facility of the Center for Transgene Technology and Gene Therapy, Campus Gasthuisberg, K.U. Leuven. Gene inactivation was obtained by homologous recombination in embryonic stem cells targeting the genes encoding tissue-type plasminogen activator (t-PA) (13), urokinase-type plasminogen activator (u-PA) (13), plasminogen activator inhibitor-1 (PAI-1) (14, 15), plasminogen (Plg) (16), or  $\alpha_2$ -antiplasmin ( $\alpha_2$ -AP) (17), as previously described. Mice with inactivated genes encoding u-PA receptor (u-PAR) (18) were not included because of the normal results obtained with u-PA deficient mice.

## 2. Materials and methods

### 2.1 Materials

Human  $\alpha_2$ -AP was prepared from fresh frozen plasma as previously described (19).

5 Polyclonal antisera were raised in rabbits by subcutaneous injection of 200 mg purified human  $\alpha_2$ -AP suspended in complete Freund's adjuvant, followed at two biweekly intervals by injection of the antigen suspended in incomplete Freund's adjuvant. Serum was obtained by  
10 repeated ear vein puncture. Pooled sera were chromatographed on Protein-A Sepharose (0.5 ml serum per ml wet gel), equilibrated with 0.1 M Tris.HCl, pH 8.1 and IgG eluted with 0.1 M glycine.HCl, pH 2.8, yielding approximately 10 mg protein per ml serum. Affino-specific  
15 antibodies were obtained from the dialyzed IgG pool by chromatography on a CNBr-activated Sepharose column substituted with human  $\alpha_2$ -AP (2.5 mg/ml wet gel) and eluted with 0.1 M glycine.HCl, pH 2.8, yielding approximately 0.1 mg specific IgG per mg applied.

20 Fab fragments were obtained from the affino-specific IgG by digestion with papain. Therefore IgG was dissolved to a concentration of 5 mg/ml and digested with 1 percent (w/w) papain in the presence of 50 mM cysteine, 1 mM EDTA, 0.1 M phosphate buffer, pH 7.0  
25 for 5 hours. The reaction was arrested by addition of iodoacetamide to a final concentration of 75 mM. After dialysis the mixture was purified on a protein A Sepharose column equilibrated with PBS. Fab concentration was determined by ELISA calibrated against an IgG  
30 standard. SDS gel electrophoresis essentially revealed homogeneous Fab fragments (not shown).

### 2.2 Production of adenoviral vectors

The recombinant adenoviruses AdCMVt-PA and  
35 AdCMVPAI-1 were generated by homologous recombination in 293 cells essentially as previously described (20). For AdCMVt-PA, an XbaI-fragment of the plasmid pSTet-PA encoding wild type human t-PA was ligated into

XbaI-digested pACCMVpLpA (21) to produce pACCMVt-PA. The adenovirus precursor pACCMVPAI-1 was generated by ligating the 1.4-kb EcoRI/BglII fragment of pPAI-1RBR containing the entire coding sequence of human PAI-1 into 5 EcoRI/BamHI-digested pACCMVpLpA. In these plasmids, the t-PA and PAI-1 cDNA are positioned between the human cytomegalovirus immediate-early enhancer/promoter and the SV40 t-antigen intron/polyadenylation signal to form a complete transcriptional unit.

10 Monolayer cultures of 293 cells (22) were cotransfected with 10 mg of pACCMVt-PA or pACCMVPAI-1 and 5 mg of pJM17 (20), a plasmid containing a full-length adenovirus 5 dl309 genome. Homologous recombination between these plasmids results in the formation of 15 recombinant viral genomes in which the adenovirus E1 region is replaced by the respective t-PA or PAI-1 transgenes. Replication of the recombinant viruses in cultured 293 cells is supported by E1A gene products supplied in trans from a copy of E1 integrated into the 20 293 cell genome.

After transfection, recombinant viral plaques were harvested and amplified as described (23). The identity of recombinant viruses was determined by restriction analysis and Southern blotting of viral DNA 25 prepared from productively infected 293 cells. The recombinant adenovirus AdRR5, which lacks an inserted gene in the E1 position, was generated from pACRR5 and pJM17 in the same manner and was used as a control adenovirus (24, 25). Recombinant viruses were replaques 30 to ensure clonal identity before further use. Large scale production of recombinant adenovirus was performed as described (23). Purified virus was supplemented with 0.1 mg/ml sterile bovine serum albumin (BSA), snap frozen in liquid nitrogen and stored at -80°C until use. The titer 35 of infectious viral particles in purified stocks was determined by plaque assay on monolayers of 293 cells with 1 hour of adsorption at 37°C. Purified viral stocks of  $>10^{10}$  plaque forming units (pfu) per ml were routinely

obtained. The kinetics and organ distribution of t-PA and PAI-1 expression following adenoviral transfer by intravenous bolus injection have been described elsewhere (26, 27).

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### 2.3 Preparation of human plasmin

Human plasminogen was prepared from fresh frozen human blood bank plasma, essentially as described previously (28). Human plasma (6 liter), to which 20 units aprotinin (Trasylol, Bayer, Germany) was added per ml, was cleared by centrifugation at 4,000 rpm for 15 min at 4°C. Lysine-Sepharose (200g wet weight, substitution level approximately 1 mg lysine per g wet Sepharose gel) was added to the supernatant, the mixture stirred for 15 hour at 4°C and the gel recovered on a Buchner funnel. Then 120 g Lysine-Sepharose was added to the filtrate, the mixture stirred and the gel recovered as above. The combined gel fractions were washed with 18 liter 0.2 M  $K_2HPO_4/KH_2PO_4$  buffer, pH 7.5, containing 10 units aprotinin per ml, then poured into a 5 x 60 cm column and washed with 0.02 M  $NaH_2PO_4$ , 0.1 M NaCl buffer, pH 7.5, containing 10 units/ml aprotinin at 4°C until the absorbance of the wash fluid at 280 nm was less than 0.05. The column was then eluted with wash buffer containing 0.05 M 6-aminohexanoic acid and protein containing fractions pooled. From 6 liter plasma approximately 145 ml fluid containing 650 mg protein was obtained. The pool was concentrated 2.5-fold on an Amicon PM10 filter and gel filtered on a 5 x 90 cm column of ultragel AcA44 equilibrated with 0.02 M  $NaH_2PO_4$ , 0.1 M NaCl buffer, pH 7.5, at a rate of 60 ml per hour. The main peak, containing approximately 590 mg protein was concentrated on an Amicon PM10 filter to a concentration of 10 mg/ml and frozen until use.

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Human plasmin was prepared from plasminogen as follows. Lysine-Sepharose (20 g wet gel) was added to human plasminogen (200 mg) solution, the mixture stirred for 3 hours at 4°C, the gel washed on a Buchner funnel



and resuspended in 30 ml 0.1 M  $\text{NaH}_2\text{PO}_4$  buffer, pH 7.4. Urokinase (500  $\mu\text{l}$  of a 50  $\mu\text{M}$  solution, prepared by activation of Saruplase (Grünenthal, Aachen, Germany) with Plasmin.Sepharose was added and the mixture stirred for 15 hours at 4°C. The gel was then washed on a Buchner funnel with 0.1 M  $\text{NaH}_2\text{PO}_4$  buffer, pH 7.4, poured into a 1.5 x 16 cm column, washed with 0.1 M  $\text{NaH}_2\text{PO}_4$  buffer, pH 7.4 until the absorbance at 280 nm of the wash fluid was less than 0.05, and eluted with 0.1 M  $\text{NaH}_2\text{PO}_4$  buffer containing 0.05 M 6-aminohexanoic acid. The protein containing fractions were pooled, glycerol was added to a final concentration of 10 percent and the pool was dialyzed at 4°C against 0.1 M  $\text{NaH}_2\text{PO}_4$  buffer containing 10 percent glycerol. The final recovery was 25 ml solution with a protein concentration of 4.0 mg/ml and an active plasmin concentration of 25  $\mu\text{M}$ .

#### 2.4 Measurement of $\alpha_2$ -antiplasmin in plasma

$\alpha_2$ -Antiplasmin levels in murine plasma were measured by a chromogenic substrate assay, based on its rapid inhibition of plasmin (29). Briefly 10  $\mu\text{l}$  mouse plasma (diluted 1/10 in 0.05 M  $\text{NaH}_2\text{PO}_4$  buffer, pH 7.4, containing 0.01% Tween 20) is mixed at 37°C with 420  $\mu\text{l}$  0.05 Tris HCl, 0.1 M NaCl buffer, pH 7.4, containing 0.01% Tween 20, and with 20  $\mu\text{l}$  of 0.125  $\mu\text{M}$  human plasmin (final concentration 5 nM). After 10s incubation, 50  $\mu\text{l}$  of 3 mM S2403 (Chromogenics, Antwerp, Belgium) is added and the change in absorbance measured at 405 nm. The change in absorbance is approximately 0.18  $\text{min}^{-1}$  with buffer and 0.09  $\text{min}^{-1}$  with pooled murine plasma.

#### 2.5 Animal experiments

Animal experiments were conducted according to the guiding principles of the American Physiological Society and the International Committee on Thrombosis and Haemostasis (30).

Focal cerebral ischemia was produced by persistent occlusion of the MCA according to Welsh et al. (31). Briefly, mice of either sex, weighing 20 to 30 g, were anesthetized by intraperitoneal injection of 5 ketamine (75 mg/ml, Apharmo, Arnhem, The Netherlands) and xylazine (5 mg/ml, Bayer, Leverkusen, Germany). Atropine (1 mg/kg; Federa, Brussels, Belgium) was administered intramuscularly, and body temperature was maintained by keeping the animals on a heating pad. A "U" shape 10 incision was made between the left ear and left eye. The top and backside segments of the temporal muscle were transsected and the skull was exposed by retraction of the temporal muscle. A small opening (1 to 2 mm diameter) was made in the region over the MCA with a hand-held 15 drill, with saline superfusion to prevent heat injury. The meningeae were removed with a forceps and the MCA was occluded by ligation with 10-0 nylon thread (Ethylon, Neuilly, France) and transsected distally to the ligation point. Finally, the temporal muscle and skin were sutured 20 back in place.

AdCMVt-PA, AdCMVPAI-1 or AdRR5 were given as an intravenous bolus of  $1.3 \times 10^9$  plaque forming units (p.f.u.) 4 days before ligation of the MCA. Human  $\alpha_2$ -AP ( $h\alpha_2$ -AP) was given intravenously divided in 2 injections, 25 given 1 min before and 30 min after ligation of the MCA, respectively. Fab fragments were injected intravenously as a bolus, 10 min after the second  $h\alpha_2$ -AP injection. Human plasmin was given intravenously as a bolus, either 15 min before or 15 min after ligation of the MCA.

30 The animals were allowed to recover and were then returned to their cages. After 24 hours, the animals were sacrificed with an overdose of Nembutal (500 mg/kg, Abbott Laboratories, North Chicago, IL) and decapitated. The brain was removed and placed in a matrix for 35 sectioning in 1 mm segments. The sections were immersed in 2% 2,3,5-triphenyltetrazolium chloride (TTC) in saline (32), incubated for 30 min at 37°C, and placed in 4 % formalin in phosphate buffered saline. With this

procedure, the necrotic infarct area remains unstained (white) and is clearly distinguishable from stained (brick red) viable tissue. The sections were photographed and subjected to planimetry. The infarct volume was defined as the sum of the unstained areas of the sections multiplied with their thickness.

The data are represented as mean  $\pm$  SEM of n determinations. The significance of differences was determined using analysis of variance followed by Fisher's PLSD test, using the Statview software package or by Student's t-test.

## EXAMPLE 2

### 15 Cerebral ischemic infarct size in mice with targeted inactivation of genes encoding plasminogen system components

Ligation of the left MCA induced a cerebral infarct with a volume of  $7.6 \pm 1.1 \text{ mm}^3$  (n= 11) in wild type mice with a mixed (50%) S129 and (50%) C57BL/6 genetic background, of  $9.3 \pm 2.7 \text{ mm}^3$  (n= 6) in inbred C57BL/6 mice and of  $6.4 \pm 1.3 \text{ mm}^3$  (n= 6) in inbred S129 mice (p= NS versus mixed background, results not shown).

Inactivation of the t-PA gene was associated with a significant reduction of infarct size to  $2.6 \pm 0.80 \text{ mm}^3$  (n= 11), (p< 0.0001 vs wild type mice), whereas inactivation of the u-PA gene had no effect on infarct size ( $7.8 \pm 1.0 \text{ mm}^3$ , n= 8, p= NS vs wild type).

Inactivation of the PAI-1 gene was associated with a significant increase in infarct size ( $16 \pm 0.52 \text{ mm}^3$ , n= 6, p< 0.0001 vs wild type) (Figure 1). In mice with inactivated Plg genes, cerebral infarct size was significantly larger than in wild type mice ( $12 \pm 1.2 \text{ mm}^3$ , n=9, p=0.037 vs wild type), whereas, conversely, in  $\alpha_2$ -AP gene deficient mice, infarct size was markedly reduced ( $2.2 \pm 1.1 \text{ mm}^3$ , n= 7, p= 0.0001 vs wild type) (Figure 1).

**EXAMPLE 3**Effect of t-PA and PAI-1 gene transfer on cerebral infarct size

Injection of  $1.3 \times 10^9$  p.f.u. of AdCMVt-PA in 6 t-PA<sup>-/-</sup> mice 4 days before MCA ligation was associated with a cerebral infarct size of  $6.0 \pm 1.3$  mm<sup>3</sup>, significantly larger than the infarcts in 5 t-PA<sup>-/-</sup> mice injected with the control virus AdRR5 ( $1.8 \pm 0.63$ ,  $p = 0.028$ ) (Figure 2A). Conversely, injection of  $1.3 \times 10^9$  p.f.u. of AdCMVP-AI-1 in 5 PAI-1<sup>-/-</sup> mice was associated with a cerebral infarct size of  $10 \pm 1.4$  mm<sup>3</sup>, significantly smaller than the infarcts in 5 PAI-1<sup>-/-</sup> mice injected with the control virus AdRR5 ( $13 \pm 1.0$  mm<sup>3</sup>,  $p = 0.019$ ) (Figure 2B).

**EXAMPLE 4**Effect of  $\alpha_2$ -antiplasmin on cerebral infarct size

Cerebral infarct size correlated with  $\alpha_2$ -AP gene dose, corresponding to  $11 \pm 2.0$ ,  $4.9 \pm 2.0$  and  $2.2 \pm 1.1$  mm<sup>3</sup> in wild type, heterozygous and homozygous deficient mice, respectively (Figure 3A). Injection of human  $\alpha_2$ -AP in groups of 4  $\alpha_2$ -AP<sup>-/-</sup> mice increased the infarct size to  $13 \pm 2.5$  mm<sup>3</sup> ( $n = 4$ ) with a 1 mg total dose and to  $11 \pm 1.5$  mm<sup>3</sup> ( $n = 6$ ) with a 0.2 mg total dose. Injection of 1.7 mg affino-specific Fab against human  $\alpha_2$ -AP in mice given 0.2 mg human  $\alpha_2$ -AP reduced the cerebral infarct size to  $5.1 \pm 1.1$  mm<sup>3</sup> ( $n = 7$ ,  $p = 0.0040$  vs 0.2 mg human  $\alpha_2$ -AP) (Figure 3B).

The above examples show that reduction of  $\alpha_2$ -AP activity (reduced  $\alpha_2$ -AP gene expression or reduction of circulating  $\alpha_2$ -AP with inhibitors) reduces focal cerebral ischemic infarct size, such as encountered during ischemic stroke.

**EXAMPLE 5**Effect of plasmin on cerebral infarct size

Injection of 50, 100 or 150  $\mu$ g human plasmin (Pli) in mice weighing approximately 30 g decreased the  $\alpha_2$ -AP levels in blood samples taken after 30 s to 67, 40 and 31 percent of baseline, respectively (mean of 2 mice, with less than 15 percent variability). Injection of 200  $\mu$ g Pli in 3 mice reduced the plasma  $\alpha_2$ -AP levels to  $59 \pm 4.8$ ,  $67 \pm 4.4$  and  $70 \pm 2.5$  percent after 2, 4 and 6 hours respectively.

Ligation of the left middle cerebral artery (MCA) induced a cerebral infarct with a volume of  $27 \pm 1.3$  mm<sup>3</sup> (n= 10) in inbred Balb/c mice, and of  $16 \pm 1.3$  mm<sup>3</sup> (n= 12) in inbred C57BL/6 mice.

Injection of 0.2 mg Pli in Balb/c mice reduced the infarct size to  $22 \pm 1.0$  mm<sup>3</sup> (n= 9) (p= 0.006 vs saline). Similar decreases were observed when the Pli injection was given 15 min before or 15 min after ligation of the MCA (Table 1). In C57Bl/6 mice, injection of 0.2 mg Pli reduced the infarct size to  $10 \pm 1.2$  mm<sup>3</sup> (n= 12) (p= 0.004 vs saline).

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\* Present address: E-317, Handa-cho 3776, Hamamatsu, Shizuoka 431-3124, Japan